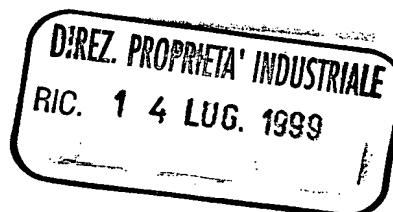


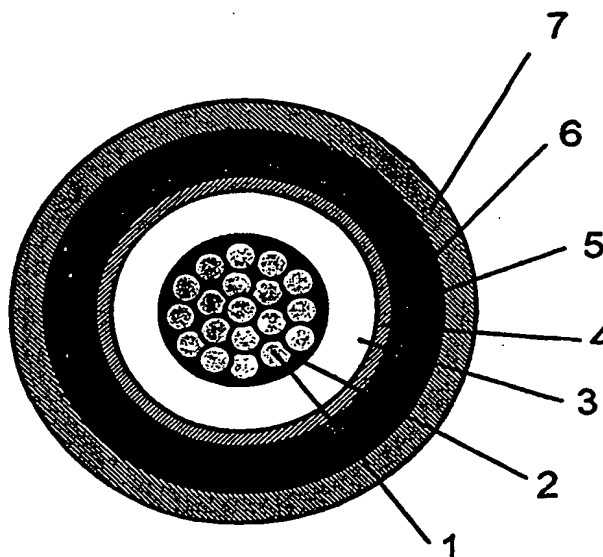
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**Published***With international search report.**Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.***(54) Title:** ELECTRICAL CABLE HAVING A SEMICONDUCTIVE WATER-BLOCKING EXPANDED LAYER**(57) Abstract**

Electrical cable, in particular for medium- or high-voltage power transmission or distribution, having a metal shield and a semiconductive water-blocking expanded layer. This layer exerts three main functions, the first one is to elastically and uniformly absorb radial forces of expansion and contraction of the cable coating layers due to thermal cycles of the cable during use, thus preventing deformations or breakages in the metal shield, the second one is to ensure electrical continuity between the cable core and the metal shield, the third one is to effectively avoid penetration and propagation of moisture and/or water along the cable core due, e.g., to possible ruptures in the metal shield. The third function is obtained by including in the expanded layer a water-swellable material.

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"ELECTRICAL CABLE HAVING A SEMICONDUCTIVE WATER-BLOCKING EXPANDED LAYER"

The present invention relates to an electrical cable,  
5 in particular for medium- or high-voltage power  
transmission or distribution, having a semiconductive  
water-blocking expanded layer. In the present description,  
the term "medium voltage" is understood to refer to a  
voltage of between about 1 kV and about 30 kV, while the  
10 term "high voltage" is understood to refer to voltages  
above 30 kV.

Cables for medium- or high-voltage power transmission  
or distribution generally consist of a metal conductor  
coated with a first inner semiconductive layer, an  
15 insulating layer and an outer semiconductive layer. For  
some uses, in particular when it needs to be watertight  
with respect to the exterior, the cable is enclosed inside  
a metal shield, usually an aluminium or copper shield,  
consisting of a continuous tube or a metal sheet shaped  
20 into a tube and welded or sealed so as to be watertight.

During production, installation or use, breakages and  
piercings can occur in the metal shield, which allow  
penetration of moisture or even water into the cable core,  
with formation of electrochemical trees in the insulation  
25 layer, which can cause insulation failure.

A possible solution to this problem is provided in US  
Patent No. 4,145,567. A high-voltage cable is described  
therein having, around the outside of the outer  
semiconductive layer, a compressible layer of a foamed  
30 plastic material which should prevent external moisture  
from reaching the insulation layer, thus avoiding  
formation of electrochemical trees. According to that  
disclosure, the metal shield preferably maintains some  
pressure against the compressible layer so that no air or  
35 other fluid can travel along the interface between the

compressible layer and the metal shield. As further insurance against passage of fluid along the cable, the metal shield can be bonded to the compressible layer. The compressible layer is preferably semiconducting.

5        Ruptures in the metal shield may be caused by the thermal cycles to which the cable is subject as a result of the daily variations in the intensity of the transported current, with corresponding variations in the cable temperature of between room temperature and the  
10        maximum operating temperature (for example between 20°C and 90°C). These thermal cycles cause dilation and subsequent contraction of the coating layers of the cable, with consequent radial forces exerted on the metal shield. The metal shield can thus suffer mechanical deformations  
15        with formation of empty spaces between the shield and the outer semiconductive layer, which may give rise to non-uniformity in the electrical field. At the utmost, these deformations can lead to rupture of the shield, particularly when it is welded or attached by means of  
20        sealing, and hence to complete loss of functionality of the shield.

      A possible solution to this problem is provided in US Patent No. 5,281,757, where the metal shield is free to move with respect to the adjacent layers and has the  
25        overlapping edge portions bonded together by an adhesive which allows the overlapping edge portions to move relative to each other during the thermal cycling of the cable. A cushioning layer as that disclosed in the above-mentioned US Patent No. 4,145,567 may be applied between  
30        the metal shield and the cable core. If desired, the cushioning layer may be a water swellable tape or a water swellable powder instead of a foamed plastic material.

      According to the Applicant's experience, cable designs such as those described in US-4,145,567 and US-  
35        5,281,757 are not completely satisfactory. Firstly, the

presence of a compressible layer between metal shield and cable core as disclosed in US-4,145,567 is not sufficient to effectively avoid penetration and propagation of moisture or water along the cable. In fact, to obtain an

5 effective water-blocking effect, in US-5,281,757 it is suggested to use, instead of the compressible layer, a water-swella-ble tape or powder. However, the introduction of a water-swella-ble material under the metal shield would cause serious electrical problems. In fact, the metal

10 shield, in addition to constituting a barrier against penetration of water and/or moisture, exerts important electrical functions and needs to be in electrical contact with the outer semiconductive layer. A first function of the metal shield is indeed to create a uniform radial

15 electric field inside the cable and, simultaneously, to cancel out the electric field outside the cable. A further function is to support short-circuit currents.

The presence of an insulating material such as a water-swella-ble material under the metal shield cannot

20 ensure electrical continuity between the cable core and the metal shield. Moreover, from the point of view of production and handling, the use of water-swella-ble tapes or of free water-swella-ble powders has many drawbacks. Particularly, the use of a water-swella-ble tape involves

25 an appreciable increase in costs and a decrease in productivity, since these tapes are expensive and imply the addition of a wrapping stage to the cable production process. On the other hand, the presence of free-flowing water-swella-ble powders makes production and installation

30 of the cable quite cumbersome.

Finally, cables are known in the art which are designed to attenuate the effect of the thermal cycles on the metal shield and at the same time to avoid propagation of moisture and/or water along the cable. These cable are

35 provided with an outer semiconductive layer with V-shaped

longitudinal grooves which are filled with a water-swella-  
ble material in the form of powder. The V-shaped  
geometry should ensure electrical contact between the  
semiconductive layer and the metal shield, on the one  
5 hand, and should assist the elastic recovery of the  
thermal dilations by the material which makes up the  
semiconductive layer, on the other hand.

However, producing these longitudinal grooves  
involves the use of a semiconductive layer of high  
10 thickness (about 2 mm or more), thereby increasing the  
cost and the overall weight of the cable. In addition, the  
desired geometry of the semiconductive layer is generally  
achieved by means of a precise process of extrusion in  
which appropriately designed dies are used. On the basis  
15 of the Applicant's experience, the formation of grooves of  
irregular geometry is, in practical terms, inevitable  
during such an extrusion process. These geometrical  
irregularities can give rise to a non-uniform distribution  
of the pressure exerted on the metal shield and thus  
20 prevent the semiconductive layer from correctly carrying  
out its function of elastic absorption of the radial  
forces.

Therefore, the cables according to the above prior  
art cannot effectively address both the problem of  
25 avoiding penetration and propagation of moisture and/or  
water inside the cable core, and the problem of possible  
deformations or breakages of the metal shield due to the  
cable thermal cycles, while maintaining a proper  
electrical contact between metal shield and cable core.

30 The Applicant has now found the above problems can be  
effectively solved by inserting, under the metal shield, a  
layer of an expanded polymer material having  
semiconductive properties and including a water-swella-  
ble material. This layer is capable of elastically and  
35 uniformly absorbing radial forces of expansion and

contraction due to the thermal cycles to which the cable is subjected during use, while ensuring the necessary electrical continuity between the cable and the metal shield. Moreover, the presence of the water-swella-  
5 material dispersed into the expanded layer is able to effectively block moisture and/or water, thus avoiding the use of water-swella- tapes or of free water-swella- powders.

In a first aspect, the present invention thus relates  
10 to an electrical cable comprising a conductor, at least one insulating layer, an outer metal shield and a layer of an expanded polymer material placed under the said metal shield, characterized in that the layer of the expanded polymer material is semiconductive and includes a water-  
15 swella- material.

In the following, the "layer of an expanded polymer material" will be concisely indicated as "expanded layer".

In the present description and claims, the term  
"expanded polymer material" is understood to refer to a  
20 polymer material with a predetermined percentage of "free" space inside the material, i.e. of space not occupied by the polymer but by a gas or air.

In general, the percentage of free space in an expanded polymer is expressed in terms of the degree of  
25 expansion (G), which is defined by the following formula:

$$G = (d_0/d_e - 1) \cdot 100$$

where  $d_0$  denotes the density of the non-expanded polymer and  $d_e$  denotes the apparent density measured on the expanded polymer.

30 The degree of expansion of the expanded layer according to the present invention can vary within a wide range, depending both on the specific polymer material used and on the thickness of the coating which it is intended to obtain. The degree of expansion is

predetermined so as to ensure that the radial forces of thermal expansion and contraction of the cable are elastically absorbed and, simultaneously, so as to maintain the semiconductive properties. In general, the  
5 degree of expansion can range from 5% to 500%, preferably from 10% to 200%.

As regards the thickness of the expanded layer according to the present invention, this is equal to at least 0.1 mm; it is preferably between 0.2 and 2 mm and  
10 even more preferably between 0.3 and 1 mm. Thicknesses below 0.1 mm are difficult to produce in practice and, in any case, only allow a limited compensation for the deformation, while thicknesses above 2 mm, although not having any functional drawbacks in principle, can be used  
15 when any specific requirements might justify the increased cost.

According to a preferred aspect, the electrical cable according to the present invention also comprises a compact semiconductive layer placed between the insulating  
20 coating and the expanded layer.

The term "compact semiconductive layer" is understood to refer to a layer made of non-expanded semiconductive material, i.e. a material having a degree of expansion substantially null.

25 According to the Applicant's perception, this compact semiconductive layer can advantageously carry out the function of preventing partial discharging, and thus damage to the cable caused by any irregularities in the surface of interface between the insulating coating and  
30 the expanded layer. This function can be performed even by a very thin semiconductive layer, namely of about 0.1 mm or even less. However, from the practical point of view, a thickness of between 0.2 and 1 mm is preferable, and even more preferably between 0.2 and 0.5 mm.

35 As indicated above, the expanded layer includes a



water swellable material. As shown by the tests carried out by the Applicant, the expanded layer is able to incorporate large amounts of water swellable material and the incorporated water-swellable material is capable of expanding when the expanded layer is placed in contact with moisture or water, thus efficiently performing its water-blocking function.

The water swellable material is generally in a subdivided form, particularly in the form of powder. The particles constituting the water-swellable powder have preferably a diameter not greater than 250  $\mu\text{m}$  and an average diameter of from 10 to 100  $\mu\text{m}$ . More preferably, the amount of particles having a diameter of from 10 to 50  $\mu\text{m}$  are at least 50% by weight with respect to the total weight of the powder.

The water-swellable material generally consists of a homopolymer or copolymer having hydrophilic groups along the polymer chain, for example: crosslinked and at least partially salified polyacrylic acid (for example the products Cabloc® from C. F. Stockhausen GmbH or Waterlock® from Grain Processing Co.); starch or derivatives thereof mixed with copolymers between acrylamide and sodium acrylate (for example products SGP Absorbent Polymer® from Henkel AG); sodium carboxymethylcellulose (for example the products Blanose® from Hercules Inc.).

To obtain an effective water-blocking action, the amount of water-swellable material to be included in the expanded layer is generally of from 5 to 120 phr, preferably of from 15 to 80 phr (phr = parts by weight with respect to 100 parts by weight of base polymer).

Figure 1 shows a cross-section of an embodiment of an electrical cable according to the present invention, of unipolar type, for medium-voltage power transmission.

This cable comprises a conductor (1), an inner semiconductive layer (2), an insulating layer (3), a

compact semiconductive layer (4), an expanded layer (5), a metal shield (6) and an outer sheath (7).

The conductor (1) generally consists of metal wires, preferably made of copper or aluminium, which are braided together using conventional techniques. The metal shield (6), usually made of aluminium or copper, or also lead, consists of a continuous metal tube or of a metal sheet shaped into a tube and welded or sealed using an adhesive material so as to make it watertight. The metal shield (6) is usually coated with an outer sheath (7) consisting of a crosslinked or non-crosslinked polymer material, for example polyvinyl chloride (PVC) or polyethylene (PE).

The polymer material which constitutes the expanded layer can be any type of expandable polymer such as, for example: polyolefins, copolymers of different olefins, copolymers of an olefin with an ethylenically unsaturated ester, polyesters, polycarbonates, polysulphones, phenol resins, urea resins, and mixtures thereof. Examples of suitable polymers are: polyethylene (PE), in particular low density PE (LDPE), medium density PE (MDPE), high density PE (HDPE), linear low density PE (LLDPE), ultra-low density polyethylene (ULDPE); polypropylene (PP); elastomeric ethylene/propylene copolymers (EPR) or ethylene/propylene/diene terpolymers (EPDM); natural rubber; butyl rubber; ethylene/vinyl ester copolymers, for example ethylene/vinyl acetate (EVA); ethylene/acrylate copolymers, in particular ethylene/methyl acrylate (EMA), ethylene/ethyl acrylate (EEA) and ethylene/butyl acrylate (EBA); ethylene/alpha-olefin thermoplastic copolymers; polystyrene; acrylonitrile/butadiene/styrene (ABS) resins; halogenated polymers, in particular polyvinyl chloride (PVC); polyurethane (PUR); polyamides; aromatic polyesters such as polyethylene terephthalate (PET) or polybutylene terephthalate (PBT); and copolymers thereof or mechanical mixtures thereof.

Preferably, the polymer material is a polyolefin polymer or copolymer based on ethylene and/or propylene, and is chosen in particular from:

5 (a) copolymers of ethylene with an ethylenically unsaturated ester, for example vinyl acetate or butyl acetate, in which the amount of unsaturated ester is generally between 5 and 80% by weight, preferably between 10 and 50% by weight;

10 (b) elastomeric copolymers of ethylene with at least one C<sub>3</sub>-C<sub>12</sub> alpha-olefin, and optionally a diene, preferably ethylene/propylene (EPR) or ethylene/propylene/diene (EPDM) copolymers, generally having the following composition: 35-90% mole of ethylene, 10-65% mole of alpha-olefin, 0-10% mole of diene (for example 1,4-  
15 hexadiene or 5-ethylidene-2-norbornene);

(c) copolymers of ethylene with at least one C<sub>4</sub>-C<sub>12</sub> alpha-olefin, preferably 1-hexene, 1-octene and the like, and optionally a diene, generally having a density of between 0.86 and 0.90 g/cm<sup>3</sup> and the following composition:  
20 75-97% by mole of ethylene; 3-25% by mole of alpha-olefin; 0-5% by mole of a diene;

(d) polypropylene modified with ethylene/C<sub>3</sub>-C<sub>12</sub> alpha-olefin copolymers, wherein the weight ratio between polypropylene and ethylene/C<sub>3</sub>-C<sub>12</sub> alpha-olefin copolymer is  
25 between 90/10 and 10/90, preferably between 80/20 and 20/80.

For example, the commercial products Elvax® (Du Pont), Levapren® (Bayer) and Lotryl® (Elf-Atochem) are in class (a), products Dutral® (Enichem) or Nordel® (Dow-Du  
30 Pont) are in class (b), products belonging to class (c) are Engage® (Dow-Du Pont) or Exact® (Exxon), while polypropylene modified with ethylene/alpha-olefin copolymers are commercially available under the brand names Moplen® or Hifax® (Montell), or also Fina-Pro®  
35 (Fina), and the like.

Within class (d), particularly preferred are thermoplastic elastomers comprising a continuous matrix of a thermoplastic polymer, e.g. polypropylene, and fine particles (generally having a diameter of the order of 1-10  $\mu\text{m}$ ) of a cured elastomeric polymer, e.g. crosslinked EPR or EPDM, dispersed in the thermoplastic matrix. The elastomeric polymer may be incorporated in the thermoplastic matrix in the uncured state and then dynamically crosslinked during processing by addition of a suitable amount of a crosslinking agent. Alternatively, the elastomeric polymer may be cured separately and then dispersed into the thermoplastic matrix in the form of fine particles. Thermoplastic elastomers of this type are described, e.g. in US-4,104,210 or EP-324,430. These thermoplastic elastomers are preferred since they proved to be particularly effective in elastically absorb radial forces during the cable thermal cycles in the whole range of working temperatures.

Products known in the art for the preparation of semiconductive polymer compositions can be used to give the polymer material semiconductive properties. In particular, an electroconductive carbon black can be used, for example electroconductive furnace black or acetylene black, and the like. The surface area of the carbon black is generally greater than 20  $\text{m}^2/\text{g}$ , usually between 40 and 500  $\text{m}^2/\text{g}$ . Advantageously, a highly conducting carbon black may be used, having a surface area of at least 900  $\text{m}^2/\text{g}$ , such as, for example, the furnace carbon black known commercially under the tradename Ketjenblack® EC (Akzo Chemie NV).

The amount of carbon black to be added to the polymer matrix can vary depending on the type of polymer and of carbon black used, the degree of expansion which it is intended to obtain, the expanding agent, etc. The amount of carbon black thus has to be such as to give the

expanded material sufficient semiconductive properties, in particular such as to obtain a volumetric resistivity value for the expanded material, at room temperature, of less than 500  $\Omega\cdot\text{m}$ , preferably less than 20  $\Omega\cdot\text{m}$ . Typically, 5 the amount of carbon black can range between 1 and 50% by weight, preferably between 3 and 30% by weight, relative to the weight of the polymer.

The compact semiconductive layer optionally present between the insulating coating and the expanded layer, as 10 well as the inner semiconductive layer, both of compact type, are prepared according to known techniques, in particular by extrusion, the polymer material and the carbon black being selected from those mentioned above for the expanded layer.

15 The insulating layer is preferably prepared by extrusion of a polyolefin selected from those mentioned above for the expanded layer, in particular polyethylene, polypropylene, ethylene/propylene copolymers, and the like. After extrusion, the material is preferably 20 crosslinked by known techniques, for example using peroxides or via silanes.

The expanded layer can be prepared by extrusion of the polymer material containing the semiconductive filler and the water-blocking material onto the core of the 25 cable, i.e. the assembly of the conductor (1), inner semiconductive layer (2), insulating layer (3) and optional compact semiconductive layer (6). The cable core can itself also be prepared by extrusion, in particular by coextrusion of the three layers according to known 30 techniques.

The polymer material can be mixed with the semiconductive filler, the water-swellaable material and other optional conventional additives according to methods known in the art. The mixing can be carried out, for

example, using an internal mixer of the type with tangential rotors (Banbury) or with interpenetrating rotors, or alternatively in continuous mixers such as those of the type Ko-Kneader (Buss), or of the type co-rotating or counter-rotating twin-screw.

The expansion of the polymer is normally carried out during the extrusion phase. This expansion can take place either chemically, by addition of a suitable expanding agent, i.e. one capable of evolving a gas under specific temperature and pressure conditions, or alternatively, physically, by high-pressure injection of gas directly into the extruder cylinder. The expanding agent is preferably added to the polymer material only after addition of the fillers and other additives as described above and subsequent cooling of the mixture below the decomposition temperature of the expanding agent to avoid a premature expansion of the polymer. Particularly, the expanding agent can be advantageously added to the polymer composition during extrusion, e.g. through the extruder hopper.

Examples of suitable expanding agents are: azodicarbamide, para-toluenesulfonyl hydrazide, mixtures of organic acids (for example citric acid) with carbonates and/or bicarbonates (for example sodium bicarbonate), and the like.

Examples of gases to be injected at high pressure into the extruder cylinder are: nitrogen, carbon dioxide, air, low-boiling hydrocarbons, for example propane or butane, halogenated hydrocarbons, for example methylene chloride, trichlorofluoromethane, 1-chloro-1,1-difluoroethane, and the like, or mixtures thereof.

Preferably, the die in the extruder head will have a diameter slightly smaller than the final diameter of the cable with expanded coating which it is intended to obtain, so that the expansion of the polymer outside the

extruder allows to reach the desired diameter for the cable.

The selected extrusion temperature mainly depends on the nature of the polymer matrix, of the expanding agent and of the desired degree of expansion. Usually, an  
5 extrusion temperature not lower than 140°C is preferred to reach a sufficient degree of expansion.

The expanded polymer material may or may not be crosslinked. The crosslinking can be carried out, after  
10 the extrusion and expansion phases, by known techniques, in particular by heating in the presence of a radical initiator, for example an organic peroxide such as dicumyl peroxide. Alternatively, a crosslinking can be carried out via silanes, which allows the use of a polymer such as  
15 those mentioned above, in particular a polyolefin, to which are covalently bonded silane units comprising at least one hydrolysable group, for example trialkoxysilane groups, in particular trimethoxysilane groups. The grafting of the silane units can take place by radical  
20 reaction with silane compounds, for example methyl triethoxysilane, dimethyldiethoxysilane, vinyl dimethoxysilane, and the like. The crosslinking is carried out in the presence of water and a crosslinking catalyst, for example an organic titanate or a metal  
25 carboxylate. Dibutyltin dilaurate (DBTL) is particularly preferred.

Once the expanded layer has been prepared, the cable is enclosed inside the metal shield. According to a preferred embodiment, in the absence of applied forces,  
30 the diameter of the expanded layer is greater than the inside diameter of the metal shield, such as to obtain, after the metal shield has been applied, a predetermined degree of precompression of the expanded layer. This precompression makes it possible to achieve optimum  
35 contact between the expanded layer and the metal shield

and can allow recovery of any residual deformation of the expanded layer, or else a certain degree of plastic deformation of the metal shield, during the phase of thermal contraction of the insulating layer.

5        Eventually, the metal shield can be coated with a protective sheath which can be obtained, for example, by extrusion of a polymer material, usually polyvinyl chloride or polyethylene.

10        A few illustrative examples will be given hereinbelow to further describe the invention.

#### EXAMPLES 1-2

15        Some mixtures suitable to form the expanded layer according to the present invention were prepared. The compositions are given in Table 1 (in phr). The components of the mixture were mixed together in a Banbury closed mixer (1.2 l working volume), loading first the base polymer, then, after a brief period of processing, the carbon black, the water swellable powder and the other additives (except the expanding agent).

20        The mixing was carried out for about 6 min with a final temperature for the extracted material of about 150°C. At the end of mixing the expanding agent was added to the mixture, the material being previously cooled to about 100°C in order to avoid a premature decomposition of the expanding agent, which would lead to uncontrolled expansion of the polymer. The mixture was then compression moulded at 160°C using a frame 200 x 200 mm in size and 3 mm in thickness. The mixture was added in amounts such as to obtain an initial layer 1 mm thick, so as to leave  
25        sufficient space for the polymer to expand. The following characteristics were measured on the test pieces thus obtained:

30        - apparent density, and then, knowing the density of the non-expanded material, degree of expansion was  
35        calculated according to the formula given above;



- volume resistivity at room temperature.

The data are given in Table 1.

Some samples were placed in water: immediate expansion of the water swellable powder up to a volume about three times the initial volume was observed.

### EXAMPLE 3

A medium-voltage cable was produced using the polymer composition of Example 1, according to the structure scheme reported in Fig. 1. The polymer composition was prepared according to Example 1, but without adding the expanding agent to avoid premature expansion of the composition. The expanding agent was introduced only during extrusion as described hereinbelow.

The cable core on which the expanded layer was to be deposited consisted of an aluminium conductor 70 mm<sup>2</sup> in cross section, coated with the following layers crosslinked via peroxide on a catenary line:

- an inner semiconductive layer made of EPR containing carbon black (0.5 mm thick);
- an insulating layer made of EPR filled with kaolin (5.5 mm thick);
- an outer semiconductive layer (compact) made of EVA containing 35% by weight of N472 carbon black (0.5 mm thick).

To deposit the expanded layer on this cable core (having an outside diameter of about 23 mm), a 80 mm single-screw extruder in configuration 25 D was used. The extruder was equipped with an initial section of the cylinder having longitudinal furrows, a box-type feed throat and a threaded discharge screw 25 D in length. The depth of the screw groove was 9.6 mm in the supply zone and 7.2 mm in the final section, for an overall screw compression ratio of about 1:1.33.

An electrically heated orthogonal extrusion head equipped with a conveyor belt with a double-suture line

was used downstream of the extruder. The following die assembly was used: tip die 24 mm in diameter, ring compression die 24 mm in diameter. The tip die was chosen with the aim of allowing easy passage of the core to be coated, with a diameter about 1 mm greater than the diameter of the core to be coated. The ring die was, on the other hand, chosen with a diameter slightly smaller than the final diameter to be obtained, so as to prevent the material from expanding inside the extrusion head.

10 The following heat profile (°C) was used for the extruder and the extrusion head:

Feed throat	Screw	Zone 1	Zone 2	Zone 3	Zone 4	Zone 5	Zone 6	Collar	Head
20	neutral	160	170	180	185	190	195	200	200

The throughput speed of the core to be coated was set as a function of the desired thickness of expanded material. In our case, a line speed of 1.2 m/min was used. Under these conditions, the following extrusion parameters were recorded:

Extruder rotation speed: 1.2 rpm;  
20 Hot semi-finished diameter: 25.0 mm;  
Cold semi-finished diameter: 24.8 mm.

The semi-finished product was cooled in air. Direct contact with the cooling water was avoided so as not to incur problems of accidental re-swelling of the water-blocking powder. The semi-finished product obtained was subsequently wound on a reel.

The material was deposited on the core in a thickness of about 1 mm. This material was expanded chemically, by adding about 2% of the expanding agent Hydrocerol® CF 70 (carboxylic acid + sodium bicarbonate) into the extruder hopper.

The electrical conductivity and the degree of

expansion were measured on samples of the expanded layer thus obtained. The degree of expansion measured was about 20%.

5 Tests of expansion of the material in the presence of water (water-blocking effect) were also carried out: the material became swollen, by virtue of the presence of the water-swellaable powder, up to a volume of about 3 times the initial volume.

#### EXAMPLE 4

10 A thermoplastic elastomer was used as base material to produce an expanded layer according to the present invention. The composition is reported in Table 1 (inclusive of the expanding agent that was added only during extrusion). The mixing was carried out in the same  
15 Banbury mixer as described for Examples 1-2 with a mixing time of about 10 min and a final temperature for the extracted material of about 195°C. After mixing, the material was granulated and sealed in plastic bags to avoid absorption of moisture.

#### 20 EXAMPLE 5

A medium-voltage cable was produced using the polymer composition of Example 4, according to the structure scheme reported in Fig. 1.

The cable core consisted of an aluminium conductor  
25 having a 150 mm<sup>2</sup> cross section and a 14.0 mm diameter, coated with the following layers, crosslinked via peroxide on a catenary line:

- an inner semiconductive layer: product LE 0595 from Borealis (0.6 mm thick);
- 30 - an insulating layer made of XLPE (4.65 mm thick);
- an outer semiconductive layer (compact): product LE 0595 from Borealis (0.4 mm thick).

The expanded layer was deposited on this core (having an outside diameter of about 25.3 mm) by extrusion  
35 according to the technique described in Example 3, using a

30 mm single-screw extruder in configuration 24 D, a tip die of 25.7 mm diameter, a ring compression die of 26.1 mm diameter, and with the following thermal profile (°C):

5

Feed throat	Screw	Zone 1	Zone 2	Zone 3	Collar	Head
20	floating	190	200	210	200	200

The expanding agent was added during extrusion through the extruder hopper. The line speed was 2.9 m/min, with a screw speed of 56 rpm. The thickness of the expanded layer after extrusion and cooling was 0.65 mm.

The so obtained cable was then wrapped with a laquered aluminium stip (thickness: 0.2 mm) using an adhesive to bond the overlapping edges. Eventually, an external sheath made of PVC was applied by extrusion.

Two 3 m portion of the final cable were subjected to a test of penetration of water under thermal cycles according to Specification NF C 33-233 March 1998). After removal of a central portion (length: 50 mm) of the external coatings to reach the outer semiconductive layer (4), the cable specimens were immersed in water and kept at room temperature for 24 hours, then subjected to 10 thermal cycles of 8 hours each (4 hours of heating up to 100°C by circulating electrical current along the conductor, then 4 hours of cooling). At the end of the test, water penetrated through the cutting for 20 cm at one side and for 25 cm at the other side, thus largely within the requirements of the specification (no water shall appear from the cable specimen extremities).

TABLE 1

EXAMPLE	1	2	4
Elvax® 470	100	--	--
Elvax® 265	--	100	--
Profax® PF 814	--	--	20
Santoprene® RC8001	--	--	80
Ketjenblack® EC 300	20	20	10
Irganox® 1010	0.5	0.5	0.2
Irganox® PS802	--	--	0.4
Waterlock® J550	40	40	25
Hydrocerol® CF70	2	2	2
$d_0$ (g/cm <sup>3</sup> )	1.15	1.15	1.012
$d_e$ (g/cm <sup>3</sup> )	0.95	0.95	0.86
Degree of expansion (%)	21	21	17.7
Volume resistivity ( $\Omega$ -m)	<15	<15	2

Elvax® 470 (Du Pont): ethylene/vinyl acetate (EVA)  
 5 copolymer (18% VA, melt index 0.7);

Elvax® 265 (Du Pont): EVA copolymer (28% VA, melt index  
 3.0);

Profax® PF 814 (Montell): isotactic propylene homopolymer  
 (MFI = 3 g/10' - ASTM D 1238);

10 Santoprene® RC8001 (Monsanto): thermoplastic elastomer  
 (89%w cured EPR, 11%w polypropylene);

Ketjenblack® EC (Akzo Chemie): high-conductivity furnace  
 carbon black;

15 Waterlock® J550 (Grain Processing Co.): grounded  
 crosslinked polyacrylic acid (partially salified) (more than 50%  
 by weight of particles having a diameter between 10 e 45  $\mu$ m);

Hydrocerol® CF70 (Boeheringer Ingelheim): carboxylic acid/sodium bicarbonate expanding agent;

Irganox® 1010 : pentaerythryl-tetrakis[3-(3,5-di-tert-butyl-4-hydroxyphenyl)propionate] (Ciba-Geigy);

5 Irganox® PS802 FL: distearyl thiodipropionate (DSTDP) (Ciba-Geigy).

CLAIMS

1. Electrical cable comprising a conductor, at least one insulating layer, an outer metal shield and a layer of an expanded polymer material placed under the said metal shield, characterized in that the layer of the expanded polymer material is semiconductive and includes a water-swella-  
5 ble material.
2. Cable according to Claim 1, wherein the expanded layer has a predetermined degree of expansion so as to ensure elastic absorption of the radial forces of thermal expansion and contraction of the cable and to maintain the semiconductive properties.  
10
3. Cable according to Claim 2, wherein the degree of expansion of the expanded layer is between 5% and 500%.  
15
4. Cable according to Claim 3, wherein the degree of expansion of the expanded layer is between 10% and 200%.  
20
5. Cable according to any one of the preceding claims, wherein the thickness of the expanded layer is at least 0.1 mm.  
25
6. Cable according to Claim 5, wherein the thickness of the expanded layer is between 0.2 and 2 mm.
7. Cable according to any one of the preceding claims, also comprising a compact semiconductive layer placed between the insulating coating and the expanded layer.  
30
8. Cable according to Claim 7, wherein the compact semiconductive layer has a thickness of from 0.1 to 1 mm.  
35

9. Cable according to Claim 8, wherein the compact semiconductive layer has a thickness of from 0.2 to 0.5 mm.

5

10. Cable according to any one of the preceding claims, wherein the water-swellaable material is in the form of powder.

10

11. Cable according to claim 10, wherein the water-swellaable material is in the form of powder having a particle diameter not greater than 250  $\mu\text{m}$  and an average particle diameter of from 10 to 100  $\mu\text{m}$ .

15

12. Cable according to claim 11, wherein in the water-swellaable material the amount of particles having a diameter of from 10 to 50  $\mu\text{m}$  are at least 50% by weight with respect the total weight of the powder.

20

13. Cable according to anyone of the previous claims, wherein the water swellaable material is a homopolymer or copolymer having hydrophilic groups along the polymer chain.

25

14. Cable according to anyone of the previous claims, wherein the water-swellaable material is present in an amount of from 5 to 120 phr.

30

15. Cable according to claim 14, wherein the water-swellaable material is present in an amount of from 15 to 80 phr.

35

16. Cable according to any one of the preceding claims, wherein the polymer material which constitutes the expanded layer is an expandable polymer selected from:



polyolefins, copolymers of different olefins, copolymers of an olefin with an ethylenically unsaturated ester, polyesters, polycarbonates, polysulfones, phenolic resins, urea resins, and mixtures thereof.

5

17. Cable according to Claim 16, wherein the polymer material is an olefin polymer or copolymer based on ethylene and/or propylene.

10 18. Cable according to Claim 17, wherein the polymer material is selected from:

(a) copolymers of ethylene with an ethylenically unsaturated ester, wherein the amount of unsaturated ester is between 5 and 80% by weight;

15 (b) elastomeric copolymers of ethylene with at least one C<sub>3</sub>-C<sub>12</sub> alpha-olefin, and optionally a diene, having the following composition: 35-90% mole of ethylene, 10-65% mole of alpha-olefin, 0-10% mole of diene;

(c) copolymers of ethylene with at least one C<sub>4</sub>-C<sub>12</sub> 20 alpha-olefin, and optionally a diene, having a density of between 0.86 and 0.90 g/cm<sup>3</sup>;

(d) polypropylene modified with ethylene/C<sub>3</sub>-C<sub>12</sub> alpha-olefin copolymers, wherein the weight ratio between polypropylene and ethylene/C<sub>3</sub>-C<sub>12</sub> alpha-olefin copolymer is 25 between 90/10 and 10/90.

19. Cable according to claim 18, wherein the polymer material is a thermoplastic elastomer comprising a continuous matrix of a thermoplastic polymer and fine 30 particles of a cured elastomeric polymer dispersed in the thermoplastic polymer.

20. Cable according to any one of the preceding claims, wherein the expanded layer has a volumetric 35 resistivity value for the expanded material at room

temperature of less than 500  $\Omega$ m.

21. Cable according to any one of the preceding claims, wherein the expanded layer comprises a  
5 predetermined amount of electroconductive carbon black.

22. Cable according to Claim 21, wherein the electroconductive carbon black has a surface area of at least 20 m<sup>2</sup>/g.

10

23. Cable according to Claim 22, wherein the carbon black has a surface area of at least 900 m<sup>2</sup>/g.

24. Cable according to any one of Claims 21 to 23,  
15 wherein the carbon black is present in amounts of between 5 and 80% by weight.

25. Cable according to Claim 24, wherein the carbon black is present in amounts of between 10 and 70% by  
20 weight.

26. Cable according to any one of the preceding claims, wherein the expanded layer is obtained by extrusion.

25

27. Cable according to Claim 26, wherein the expansion of the layer is obtained during the extrusion by addition of an expanding agent.

30 28. Cable according to Claim 26, wherein the expansion of the layer is obtained during extrusion by high-pressure injection of a gas.

29. Cable according to any one of the preceding

claims, wherein the diameter of the expanded layer, in the absence of applied forces, is greater than the inside diameter of the metal shield, so as to obtain a predetermined degree of precompression of the expanded layer after the metal shield has been applied.

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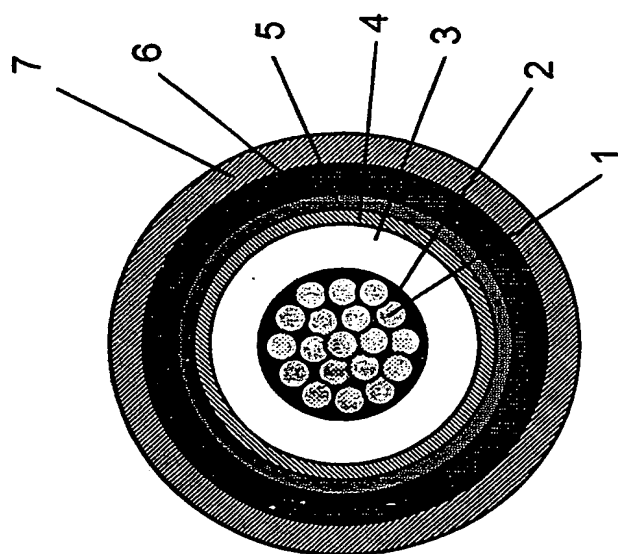


Fig. 1

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## INTERNATIONAL SEARCH REPORT

Internatio application No  
PCT/EP 8284A. CLASSIFICATION OF SUBJECT MATTER  
IPC 6 H01B9/02

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 H01B

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 4 145 567 A (BAHDER ET AL.) 20 March 1979 see column 2, line 35 - column 5, line 68; figure 1	1,2,7, 16-18
A	EP 0 116 754 A (CABLE TECHNOLOGY LABORATORIES) 29 August 1984 see page 7, line 34 - page 13, line 4; figure 1	1,2,7
A	EP 0 577 233 A (SIEMENS) 5 January 1994 see column 2, line 1 - line 27; figures 1,2	1,7,10

☐ Further documents are listed in the continuation of box C.☒ Patent family members are listed in annex.

## \* Special categories of cited documents :

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# INTERNATIONAL SEARCH REPORT

Information on patent family members

Automatic Application No  
PCT/EP 98/08284

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